The hospital anxiety and depression scale, in patients with multiple sclerosis

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Purpose: Multiple sclerosis (MS) is a long-lasting disabling disease with psychological implications. Hospital Anxiety and Depression Scale (HADS) is a questionnaire developed to screen anxiety and depression among patients in hospital settings. The objective of this report was to study the metric properties of the Portuguese version of HADS regarding MS.

Materials and methods: This was a cross-sectional study. Three hundred and eighty-five individuals with a diagnosis of MS, according to the revised McDonald criteria, were recruited from an outpatient Neuroimmunology Clinic of the city of Porto. Participants had a mean age of 40.04 years (SD =10.86), 63.9% of them were female. HADS consists of two subscales – anxiety and depression – scored separately, with seven items each.

Results: The results of our study show the appropriateness of HADS for the identification of mood changes in people with MS and show good metric properties.

Conclusion: It seems to be a useful tool for the scrutiny of mood disorders in Portuguese patients with MS. The questionnaire is also sensitive to disease when compared to people with other severe illnesses, and with people without the disease.

Keywords: multiple sclerosis, depression, anxiety, HADS

Introduction

Multiple sclerosis (MS) is a chronic disease of the nervous system that produces various symptoms and may lead to severe disability with critical psychological implications. Population-based studies show that most patients are ambulatory even after many years of disease duration.1

The Hospital Anxiety and Depression Scale (HADS) is a brief self-reporting two-dimensional questionnaire developed to screen for levels of anxiety and depression among patients in a general hospital setting.2,3 In a systematic review of the structure of the HADS, Cosco et al4 found divergences in the latent structure of the questionnaire and recommends caution because of the apparent lack of consistency between studies. This matter led to calls for the abandonment of the HADS.5 In response to these conclusions, a meta confirmatory factor analysis (CFA)6 provides support for the bifactor model but suggests caution in the use of the scale. Stern7 confirms that HADS is appropriate for initial diagnosis and to track the progression of psychological symptoms. National Institute for Health and Care Excellence (section 1.3.2, point 1.3.2.3)8 had recommended the scale for the diagnosis of depression and anxiety.

HADS is one of the most widely used questionnaires among patients with neurological illnesses, and it is useful for assessing patients who tire easily or have limited concentration spans.9 Watson et al10 confirm that HADS is an appropriate screening measure for people with MS. Litster et al11 in a systematic review about the assessment...
of anxiety in people with MS state that HADS-A shows promise as an applicable measure for this population. In the evaluation of depression in people with MS, a systematic review of Hind et al12 asserts that HADS-D is a useful screening instrument.

However, research alerts about possible problems in the translated versions and cross-cultural use of the HADS and claims that rates of anxiety and depression using the same cut point across languages and culture may be wrong or subject to critical scrutiny.13

In a previous study,14 the metric properties of a Portuguese version of HADS were described when applied to 1,322 individuals with chronic diseases, namely, cancer, stroke, epilepsy, coronary heart disease, morbid obesity, diabetes, myotonic dystrophy, obstructive sleep apnea, depression, and a non-disease group.

The core objective of the present report was to describe the metric properties of the Portuguese version of HADS among people with MS.

Materials and methods

Participants

We analyzed a consecutive sample of 380 individuals (63.9% female, 36.1% male; mean age 40.04±10.86 years; range: 16–71 years) with MS according to the revised McDonald criteria, from the outpatient Neuroimmunology Clinic at a Central Hospital in Porto, Portugal.

Measure

HADS

HADS consists of two subscales, one measuring anxiety, with seven items, and another measuring depression, with seven items, which score separately. Each item was answered by the patient on a 4-point (0–3) scale, so the possible scores ranged from 0 to 21 for each of the two subscales, taking 2–5 minutes to complete.2,3 The HADS manual indicates that a score between 0 and 7 is “normal,” between 8 and 10 “mild,” between 11 and 14 “moderate,” and between 15 and 21 “severe.” Some authors propose a different classification. In a study with a Brazilian sample (Portuguese-Brazilian language), Botega et al15 proposed a cutoff point of 8/9 for anxiety and depression scales. Other studies suggest that the recommended cutoff scores for the HADS may result in under-reporting of psychiatric morbidity in different chronic diseases.16

The original authors post a score of ≥11 as indicating the probable presence (“caseness”) of a mood disorder, and a score of 8–10 being just suggestive of the presence of the respective state. Based on the score of ≥8, Honarmand and Feinstein’s research17 confirms the usefulness of the HADS as a marker of major depression and generalized anxiety disorder in MS patients. In a review of different screening measures to identify anxiety disorders in MS, Litster et al11 conclude that HADS is the best available tool for detecting the symptoms of anxiety in individuals with MS.

Procedure

The study used a cross-sectional design with all measures taken at the same time. Ethical approval for the study was given by the ethical committee at the hospital St. Antonio, Porto, according to hospital rules and Portuguese law. Namely, signed informed consent was obtained before participation in the study in accordance with the Declaration of Helsinki. A registered psychologist administered the tests individually during the patient routine visit to the hospital.

Statistical analysis

To identify the factorial structure of the scale, in this group of MS patients, exploratory factor analysis was applied. Moreover, to test the hypothesized two model dimensions, CFA was used.

Results

Exploratory factorial validity

Exploratory factor analysis using principal component analysis, oblimin rotation method with Kaiser normalization for the 14 items, showed two components: the first one associated with the seven anxiety items and the second one related to the seven items from depression. The total variance of the solution explains 49.07% (39.26% for the first component). Loadings on the component that they belong to are in bold format in Table 1. All items load on the dimension they belong to.

Differences in loadings between the component the item belongs to and the component it does not belong to are above 0.15 points, except for item 14. Table 1 shows loadings on the dimensions above 0.35. Internal consistency, using Cronbach’s alpha, for the two dimensions is good; 0.80 for anxiety and 0.81 for depression.

CFA

CFA is a theory-testing model as opposed to a theory-generating method such as exploratory factor analysis. Byrne19 suggests that there is a wide array of fit indexes from which to choose, but that only one or two needs be reported. Maximum likelihood CFA, using EQS V6.121 was used to test the two-factor
Table 1 Exploratory factor analysis – presenting loadings above 0.35

<table>
<thead>
<tr>
<th></th>
<th>Anxiety</th>
<th>Depression</th>
</tr>
</thead>
<tbody>
<tr>
<td>HADS1</td>
<td>0.74</td>
<td>0.39</td>
</tr>
<tr>
<td>HADS2</td>
<td>0.78</td>
<td>0.39</td>
</tr>
<tr>
<td>HADS3</td>
<td>0.75</td>
<td>0.40</td>
</tr>
<tr>
<td>HADS4</td>
<td>0.49</td>
<td>0.79</td>
</tr>
<tr>
<td>HADS5</td>
<td>0.74</td>
<td>0.40</td>
</tr>
<tr>
<td>HADS6</td>
<td>0.53</td>
<td>0.79</td>
</tr>
<tr>
<td>HADS7</td>
<td>0.60</td>
<td>0.43</td>
</tr>
<tr>
<td>HADS8</td>
<td>0.53</td>
<td>0.79</td>
</tr>
<tr>
<td>HADS9</td>
<td>0.65</td>
<td>0.43</td>
</tr>
<tr>
<td>HADS10</td>
<td>0.35</td>
<td>0.72</td>
</tr>
<tr>
<td>HADS11</td>
<td>0.52</td>
<td>0.71</td>
</tr>
<tr>
<td>HADS12</td>
<td>0.36</td>
<td>0.71</td>
</tr>
<tr>
<td>HADS13</td>
<td>0.71</td>
<td>0.44</td>
</tr>
<tr>
<td>HADS14</td>
<td>0.46</td>
<td>0.50</td>
</tr>
</tbody>
</table>

Notes: Extraction method: principal component analysis. Rotation method: oblimin with Kaiser normalization. Bold figures represent that the item belongs to that factor or dimension in coincidence with the higher loading.

Table 2 Anxiety and depression means by disease group, and statistically significant differences with MS patients

<table>
<thead>
<tr>
<th>Disease group</th>
<th>n</th>
<th>Anxiety</th>
<th>Depression</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>M (SD)</td>
<td>t</td>
</tr>
<tr>
<td>No disease</td>
<td>22</td>
<td>7.81 (4.72)</td>
<td>0.09</td>
</tr>
<tr>
<td>Cancer</td>
<td>98</td>
<td>9.18 (4.74)</td>
<td>2.59</td>
</tr>
<tr>
<td>Stroke</td>
<td>252</td>
<td>10.53 (4.36)</td>
<td>7.50</td>
</tr>
<tr>
<td>Epilepsy</td>
<td>100</td>
<td>8.59 (4.17)</td>
<td>1.45</td>
</tr>
<tr>
<td>Diabetes</td>
<td>316</td>
<td>8.18 (4.53)</td>
<td>0.86</td>
</tr>
<tr>
<td>Cardiac coronary disease</td>
<td>274</td>
<td>8.41 (3.86)</td>
<td>1.57</td>
</tr>
<tr>
<td>Morbid obesity</td>
<td>190</td>
<td>10.52 (4.35)</td>
<td>6.85</td>
</tr>
<tr>
<td>Obstructive sleep apnea</td>
<td>31</td>
<td>6.87 (3.39)</td>
<td>1.31</td>
</tr>
<tr>
<td>Depression</td>
<td>20</td>
<td>11.00 (3.91)</td>
<td>3.18</td>
</tr>
<tr>
<td>Myotonic dystrophy</td>
<td>18</td>
<td>7.27 (4.07)</td>
<td>0.61</td>
</tr>
<tr>
<td>Multiple sclerosis</td>
<td>380</td>
<td>7.94 (4.31)</td>
<td>0.46</td>
</tr>
</tbody>
</table>

Notes: HADS, Hospital Anxiety and Depression Scale; MS, multiple sclerosis; SD, standard deviation; P, level of statistical significance; t, test for independent samples.

For depression by gender, 9.9% of females exhibit a value of ≥11 and 28.8% a value of ≥8, and for anxiety, 21.6% of the participants presented a score of ≥11 and 41.6% a score of ≥8. These values are similar to those reported by Siegert and Abernethy. For depression, report higher values in men: these differences are also reported in other studies in different languages and problems. In the present study, for depression, 8.2% of the participants exhibited a score of ≥11 and 23.4% a score of ≥8, and for anxiety, 21.6% of the participants presented a score of ≥11 and 41.6% a score of ≥8. These values are similar to those reported by Jones et al. Differences between gender for anxiety and depression are statistically significant at the P level <0.001, with females showing higher mean values than males (anxiety 8.72 vs 6.57 and depression 6.24 vs 4.55). These values are different from the ones reported by Jones et al which, for depression, report higher values in men: these differences are also reported in other studies in different languages and problems. However, correlations between age and depression is statistically significant [r(375)=0.30, P<0.001], also different from the study of Jones et al which did not find statistically significant correlations between age and depression and report a statistically significant correlation between age and anxiety.

When comparing the MS group with other chronic diseases, more statistically significant differences are found for depression than for anxiety. For anxiety, no statistically significant correlations between age and anxiety are not statistically significant. However, correlations between age and depression is statistically significant [r(375)=0.30, P<0.001], also different from the study of Jones et al which did not find statistically significant correlations between age and depression and report a statistically significant correlation between age and anxiety.
significant differences were found between the MS group and the non-disease group. For depression, statistically significant differences \(t(390) = 2.74, P < 0.01\) were present, showing that the MS group had higher values \((M = 5.63)\) than the non-disease group \((M = 3.22)\). Comparing with the depression disease group, the magnitude of values is much higher for the depression group. These results seem to confirm the results of a review by Siegert and Abernethy,\(^{29}\) which states that depression is common in MS.

Statistically significant differences were found when anxiety and depression scores were compared between MS patients and those with other diseases (Table 2). Compared with the depression group, MS patients showed statistically significant lower values in anxiety \(t(388) = 3.18, P < 0.01\) and in depression \(t(388) = 7.87, P < 0.001\).

Furthermore, similar differences were found in anxiety scores in cancer, stroke, and morbid obesity patients, and in depression scores in stroke and morbid obesity patients. MS patients had significantly higher depression scores than obstructive sleep apnea patients.

Correlations between age and anxiety and depression show a non-significant association for anxiety and a statistically significant association – though moderate – for depression \(r(365) = 0.29, P < 0.0001\), suggesting that age is not associated with anxiety and depression. The correlation between anxiety and depression \(r(380) = 0.60, P < 0.0001\) is similar to the Portuguese adaptation study\(^{14,19}\) and other reported studies.

**Discussion and conclusion**

International research on HADS shows that it is a useful tool to evaluate anxiety and depression in people suffering from different diseases, including MS. In an extensive study, Dahl et al\(^{15}\) show that anxiety and depression occur more frequently in MS patients than in the general population. Because anxiety and depression seem to be common in people with MS,\(^{17}\) it seems appropriate to screen for possible mood elevation in these patients. Our study identifies, for depression, essential differences between people with MS and people without diseases, and people with a diagnostic of depression and people with MS, suggesting the existence of a sensibility that differentiates between people with MS and people with and without the diagnosis of depression when using HADS dimensions.

The results of our report confirm the findings from international research and suggest that the Portuguese HADS version behaves similarly. It seems to be a useful tool for the scrutiny of mood in Portuguese patients with MS as it is for other diseases. Remember that in the original development of the scale, because scores can be affected by the physical illness of the patient, authors eliminated items relating simultaneously to emotional disorder and physical disorder. Items selected were based solely on the anxiety and depression symptoms.\(^3\) Because items are independent of the characteristics of the disease, people with different diseases, and people without diseases, may have the same level of emotional disturbance, due to some other life events than the ones related to the disease (symptoms and inherent treatment). HADS is a useful tool to help people in medical settings cope with the disease and reduce emotional distress.

The objective of the research was to report the metric properties and the dimensionality of HADS because recent research suggests for example “No further research needed: abandoning the Hospital and Anxiety Depression Scale.”\(^{15}\) While others suggests a different number of dimensions of the scale.\(^{14,31,32}\) However, more profound research recommends that the two-factor solution seems to be the most appropriate solution for HADS\(^6\) together with large population studies.\(^{18}\) Other research suggests that HADS may be challenging to translate and that translated versions of the HADS may not be attainable.\(^{19}\) However, different languages found that the HADS is appropriate for use and found the same factorial solution for the HADS.\(^{14,33}\) In the present study, the results suggest that the translated version behaves like other different languages translations. The psychometric properties are good, and a clinimetric analysis shows sensitivity when we compare people with different chronic diseases to people without any condition. In conclusion, the HADS seems to be a useful instrument to be included in the study of people with MS.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**


